mL of CH_2Cl_2, and the combined organics were dried over K_2CO_3 and concentrated *in vacuo*. Chromatography on 75 g of silica gel, eluting with 35% acetone/hexanes + 1% Et_3N afforded 218 mg (27%) of **8b**: ^{1}H NMR δ 1.35 (s, 9H), 3.59 (bs, 1H), 7.02–7.08 (m, 2H), 7.97–7.99 (m, 1H); ^{13}C NMR δ 29.9, 51.5, 122.7, 123.3, 139.3, 143.0; R_f 0.42, 50% acetone/hexanes + 1% Et_3N; IR 3396, 1584, 1482, 1227 cm^{-1}; HRMS calcd for $C_9H_{14}N_2$ 151.1235, found 151.1223.

Supplementary Material Available: Copies of ^{1}H NMR spectra of **2**, **3**, **5**, **7a–c**, **7e**,**f**, **8a**,**b** (10 pages). This material is contained in libraries on microfiche, immediately follows this article in the microfilm version of the journal, and can be ordered from the ACS; see any current masthead page for ordering information.

JO950036N

Additions and Corrections

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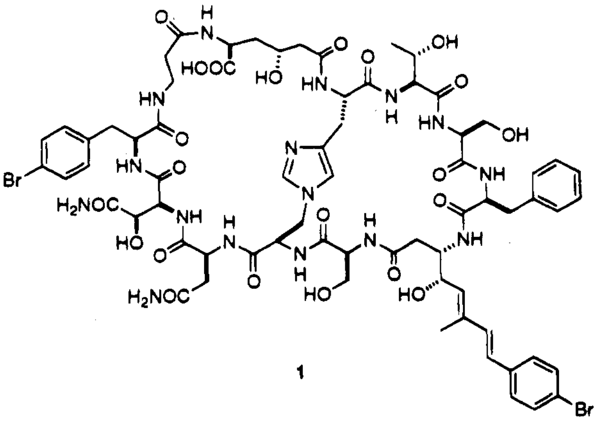
Louis D. Quin* and Stefan Jankowski. *O*-Ethyl Phosphoramidic Acids with Sterically Demanding *N*-Substituents: Useful Precursors of Ethyl Metaphosphate on Thermolysis.

Page 4408, column 1, line 3 of experimental section, replace Et_2NH with Et_3N. Line 6, replace $Et_2NH{\cdot}HCl$ with $Et_3N{\cdot}HCl$. Following line 9, replace one OH of the phosphoramidic acid precursor of **17** by OEt.

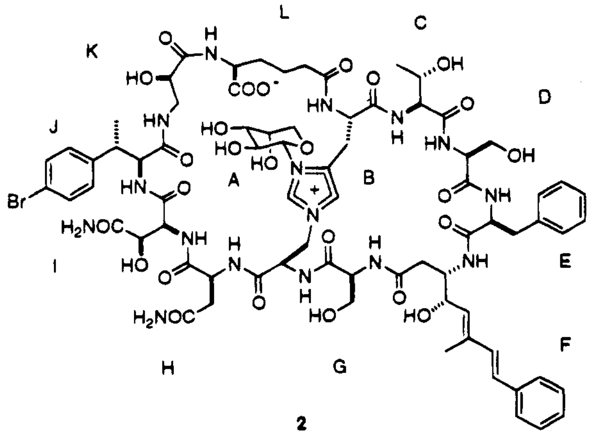
JO944015P

Carole A. Bewley and D. John Faulkner*. Theonegramide, an Antifungal Glycopeptide from the Philippine Lithistid Sponge *Theonella swinhoei*.

Page 4849, Chart 1. The methyl group in the AHMP residue (F) in theonegramide (**2**) should be at C-6. Theonellamide F (**1**) was also drawn incorrectly.



1



2

JO954002R

James A. Marshall* and Chad E. Bennett. Synthesis of Furans by S_N2' Cyclization of γ-Alkynyl Allylic Alcohol Derivatives.

Page 6133. Incorrect numbers were assigned to compounds **13a**, **13b**, and **14** in the Experimental Section of this paper. The corrected version should read as follows.

(*Z*)-6-Hydroxy-1-hexyl-1,4-dimethyl-4-hexen-2-ynyl 2,6-Dimethoxybenzoate (13b). The procedure described for benzoate **4b** was followed with 2.00 g (5.91 mmol) of (*Z*)-1-((*tert*-butyldimethylsilyl)oxy)-3,6-dimethyl-2-dodecen-4-yn-6-ol[1] and 3.56 g (17.7 mmol) of 2,6-dimethoxybenzoyl chloride for 47 h. The product was purified by flash chromatography on silica gel (10%EtOAc–hexane, followed by 25%EtOAc–hexane) to yield 2.64 g (89%) of benzoate as a yellow oil: ^{1}H NMR (300 MHz, $CDCl_3$) δ 7.23 (t, J = 8.4 Hz), 6.51 (d, J = 8.4 Hz), 5.73 (dt, J = 1.5, 4.9 Hz), 4.38 (dt, J = 1.3, 5.1 Hz), 3.78 (s), 1.85 (q, J = 1.3, 1.4 Hz), 1.79 (s), 1.52–1.28 (m), 0.87 (m), 0.05 (m).

The procedure described for alcohol **5b** was followed with 2.64 g (5.25 mmol) of the foregoing benzoate for 45 h. The product was purified by flash chromatography on silica gel (25% EtOAc–hexane, followed by 50% EtOAc–hexane) to yield 0.541 g (27%) of a pure benzoate **13b** and 1.190 g (58%) of slightly impure benzoate **13b**: ^{1}H NMR (300 MHz, $CDCl_3$) δ 7.24 (t, J = 8.4 Hz), 6.52 (d, J = 8.4 Hz), 5.91 (dt, J = 1.5, 5.3 Hz), 4.29 (d, J = 6.9 Hz), 3.79 (s), 2.01–1.92 (m), 1.88 (d, J = 1.5 Hz), 1.77 (s), 1.70 (s), 1.53–1.28 (m), 0.87 (t, J = 6.7 Hz). Anal. Calcd for $C_{23}H_{32}O_5$: C, 71.11; H, 8.30. Found: C, 70.86; H, 8.32.

(*E*)-2-(2-Methyl-1-octenyl)-3-methylfuran (14): A. Cyclization of MOM Ether 13a with KO-t-Bu in THF. Procedure A described for furan **6** was followed with 0.255 g (0.950 mmol) of MOM ether **13a** for 3.75 h. The product was purified by flash chromatography on silica gel (2.5% EtOAc–hexane) to yield 0.164 g (84%) of furan **14** as a 1:1 mixture of *E* and *Z* isomers: ^{1}H NMR (300 MHz, $CDCl_3$) δ 7.27 (d, J = 1.8 Hz), 7.25 (s), 6.19 (d, J = 2.6 Hz), 6.18 (d, J = 2.0 Hz), 5.92 (s), 5.90 (s), 2.43 (t, J = 7.5 Hz), 2.12 (t, J = 7.5 Hz), 1.99 (s), 1.86 (d, J = 1.3 Hz), 1.46–1.28 (m), 0.86 (m).

B. Cyclization of 2,6-Dimethoxybenzoate 13b with KOH–Aliquot 336. Procedure D described for furan **6** was followed with 0.196 g (0.504 mmol) of benzoate **13b** at 55–65 °C for 1.75 h. The product was purified by flash chromatography on silica gel (2.5% EtOAc–hexane) to yield 0.081 g (78%) of furan **14**, a light yellow oil, as a 60:40 mixture of *E* and *Z* isomers.

JO954003J